		<b>AETNA BETTER HEALTH®</b> Coverage Policy/Guideline	
Name:	Eucrisa	Page:	1 of 2
Effective Date:	2/3/2025	Last Review Date:	12/17/2024
Applies to:	<input checked="" type="checkbox"/> Illinois <input checked="" type="checkbox"/> Maryland	<input checked="" type="checkbox"/> Florida Kids <input checked="" type="checkbox"/> Pennsylvania Kids	<input checked="" type="checkbox"/> New Jersey <input type="checkbox"/> Virginia

### Intent:

The intent of this policy/guideline is to provide information to the prescribing practitioner outlining the coverage criteria for Eucrisa under the patient's prescription drug benefit.

### Description:

Eucrisa is indicated for topical treatment of mild to moderate atopic dermatitis in adult and pediatric patients 3 months of age and older.

### Applicable Drug List:

Preferred Agent: Eucrisa

### Policy/Guideline:

The requested drug will be covered with prior authorization when the following criteria are met:

- The requested drug is being prescribed for mild to moderate atopic dermatitis in a patient 3 months of age or older

#### AND

- The patient is less than 2 years of age

#### OR

- The requested drug will be used on sensitive skin areas (e.g., face, genitals, or skin folds)

#### AND

- The patient experienced an inadequate treatment response, intolerance, or contraindication to a topical calcineurin inhibitor

#### OR


- The patient experienced an inadequate treatment response, intolerance, or contraindication to a topical calcineurin inhibitor AND a medium or higher potency topical corticosteroid

#### AND

- If additional quantities are being requested, then 5 percent or greater body surface area is affected

#### OR

- The request is for continuation of therapy, and the patient achieved or maintained positive clinical response as evidenced by improvement [(e.g., improvement in or resolution of any of the following signs and symptoms: erythema (redness), exudation (oozing and crusting), excoriation (evidence of scratching), induration (hardening)/papulation (formation of papules), lichenification (epidermal thickening), OR pruritus (itching)]

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**AND**

- If additional quantities are being requested, then 5 percent or greater body surface area is affected

**Approval Duration and Quantity Restrictions:**

**Approval:**

- Initial – 3 months
- Renewals – 12 months

**Quantity Level Limit:**

- 60gm/30 days
- 120gm/30 days when 5% or greater of body surface area (BSA) is affected

**References:**

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4. Eichenfield LF, Tom WL, et. al. Guidelines of care for the management of atopic dermatitis: Section 2. Management and treatment of atopic dermatitis with topical therapies. *J Am Acad Dermatol*. 2014;71:116-32.
5. Paller AS, Tom WL, et. al. Efficacy and safety of crisaborole ointment, a novel, nonsteroidal phosphodiesterase 4 (PDE4) inhibitor for the topical treatment of atopic dermatitis (AD) in children and adults. *J Am Acad Dermatol*. 2016 Jul 1175(3)494-503.e4.
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7. Eichenfield LF, Tom WL, et. al. Guidelines of Care for the Management of Atopic Dermatitis: Section 1. Diagnosis and assessment of atopic dermatitis. *J Am Acad Dermatol* 2014; 70:338-51.
8. Sidbury RS, Alikhan A, Berovitch L, et al. Guidelines of care for the management of atopic dermatitis in adults with topical therapies. *J Am Acad Dermatol*. 2023; 89(1): e1-e20.